N S AD A 0 604 rept. Oct 77- Sep 78 CE1-711-1036 TN no7 N-1526 PERFORMANCE OF LEAD-ACID GELLED ELECTROLYTE BATTERIES IN THE DEEP OCEAN ENVIRONMENT, author: W. D./Briggs date:/sep.man.978 sponsor: NAVAL MATERIAL COMMAND Washington, DC 20360 program nos: ZF61 512 001 079 X F61512-1 ENGINEERING LABORATORY NAVAL CONSTRUCTION BATTALION CENTER Port Hueneme, California 93043 Approved for public release; distribution unlimited.

70 m

nit

Unclassified

SECURITY CLASSIFICATION OF THIS PAGE (When Data Entered)

·		
REPORT DOCUMENTATION	PAGE	READ INSTRUCTIONS BEFORE COMPLETING FORM
I REPORT NUMBER	2 GOVT ACCESSION NO	
TN-1526 /	DN887003	
4. YITLE (and Subtitle)		5 TYPE OF REPORT & PERIOD COVERED
PERFORMANCE OF LEAD-ACID GEL		Final; Oct 1977 - Sep 1978
ELECTROLYTE BATTERIES IN THE	DEEP OCEAN	6 PERFORMING ORG. REPORT NUMBER
ENVIRONMENT		
7 AUTHOR(s)		B CONTRACT OR G ANT NUMBER(s)
W. D. Briggs		
9. PERFORMING ORGANIZATION NAME AND ADDRESS CIVIL ENGINEERING LABORATORY	. /	10 PROGRAM ELEMENT, PROJECT, TASK AREA & WORK UNIT NUMBERS
Naval Construction Battalion Center		62766N;
Port Hueneme, California 93043		ZF61-512-001-079
11. CONTROLLING OFFICE NAME AND ADDRESS		12 REPORT DATE
Naval Material Command		September 1978
Washington, DC 20360		13 NUMBER OF PAGES
14 MONITORING AGENCY NAME & ADDRESS(It different	t from Contexting Office)	24 15 SECURITY CLASS (of this report)
The month of the desired that a noncessia different	mon contouring office)	13 SECONTY CERSS (of ans report)
		Unclassified
		15a DECLASSIFICATION DOWNGRADING SCHEDULE
16 DISTRIBUTION STATEMENT (of this Report)		
17 DISTRIBUTION STATEM ANT (of the abstract entered in	in Block 20, if different fro	m Report)
18 SUPPLEMENTARY NOTES		
19 KEY WORDS (Continue on reverse side if necessar) and	d identify by block numbe	
Batteries, gelled electrolyte, power source	es, deep ocean powe	er sources.
20 ABSTRACT (Continue on reverse side If necessary and		
> Standard lead-acid batteries with lic when charged and discharged under deep lead-acid batteries with gelled electrolyte pressure and low temperature.	ocean conditions. were tested to dete	Recently developed commercial rmine their performance at high
Batteries from three manufacturers	were divided into o	control and test groups. The
		continued

DD 1 FORM 1473 EDITION OF 1 NOV 65 IS OBSOLETE

Unclassified
SECURITY CLASSIFICATION OF THIS PAGE (When Data Entered)

SECURITY CLASSIFICATION OF THIS PAGE (When Date Entered)

20. Continued

test groups were modified to allow pressure compensation with white mineral oil. The performance of the test batteries was not degraded by exposure to the mineral oil or high pressure (10,000 psig). An average loss of capacity of 10% was experienced when the batteries were cycled cold (32°F) and at pressure (10,000 psig). However, this capacity loss is significantly less than the 35-70% loss experienced in previous tests with standard batteries in similar conditions. Thus, lead-acid batteries with gelled electrolyte were found to be suitable for use in deep ocean conditions in a pressure compensated mode.

Library Card

Civil Engineering Laboratory

PERFORMANCE OF LEAD-ACID GELLED ELECTROLYTE BATTERIES IN THE DEEP OCEAN ENVIRONMENT (Final),

by W. D. Briggs

TN-1526 24 pp illus

September 1978

Unclassified

1. Batteries

2. Gelled electrolyte

I. ZF61-512-001-079

Standard lead-acid batteries with liquid electrolyte experience dramatic capacity losses when charged and discharged under deep ocean conditions. Recently developed commercial lead-acid batteries with gelled electrolyte were tested to determine their performance at high pressure and low temperature.

Batteries from three manufacturers were divided into control and test groups. The test groups were modified to allow pressure compensation with white mineral oil. The performance of the test batteries was not degraded by exposure to the mineral oil or high pressure (10,000 psig). An average loss of capacity of 10% was experienced when the batteries were cycled cold (32°F) and at pressure (10,000 psig). However, this capacity loss is significantly less than the 35-70% loss experienced in previous tests with standard batteries in similar conditions. Thus, lead-acid batteries with gelled electrolyte were found to be suitable for use in deep ocean conditions in a pressure compensated mode.

Unclassified

SECURITY CLASSIFICATION OF THIS PAGE(When Data Entered)

Table of Contents

	Pa	<u>age</u>
INTRODUCTION	•	1
TEST PROCEDURE	•	1
TEST RESULTS	•	3
CONCLUSIONS	, .	6
ACKNOWLEDGEMENTS	• •	7
REFERENCE	•	7
APPENDIX A	• •	14
APPENDIX B		15

ACCESSION NTIS	IV., ta Section (1)
DDC	E if Section 🗍
UNANTIOUT!	ore D
JEST 107 (l' i
ov Disable '	ज्यात १११ - में ४ ५५ दिनाई
1 815 1 2	ان <u>ب</u>
1	1

INTRODUCTION

Lead-acid batteries have traditionally been the major source of electrical power for deep ocean applications. They are readily available, reasonably priced and reliable in most situations. However, tests and experience with standard cells containing liquid electrolyte (H2SO4) have shown a marked degradation in performance when the ambient environment is both cold (near 0°C) and hyperbaric. It has been hypothesized that the electrolyte stratifies at low temperature because the high pressure eliminates the hydrogen gas bubbles which normally rise from the plates thereby forcing circulation of the electrolyte. Previous unpublished test results showed capacity losses of about 35% at 6000 psi and temperatures from 32-35°F. This capacity loss increased to as much as 70% when the batteries were allowed to stand in these deep ocean conditions for five days prior to discharge.

Recent commercial developments have allowed the production of leadacid cells with the electrolyte in a gelled or paste form. Because of the gelled form of the electrolyte, it was believed that the electrolyte would not stratify, thus improving deep ocean lead-acid battery performance. The purpose of these tests was to establish the suitability and performance of gelled electrolyte batteries in a pressure compensated mode exposed to deep ocean conditions. If these batteries will perform in the test environment without experiencing major capacity losses, a reliabile power source that would not have to be contained within a pressure housing would be commercially available.

TEST PROCEDURE

To test the theory that gelled electrolyte lead-acid batteries could produce improved results for many deep ocean applications, six 6-volt batteries were procured from 3 commercial suppliers as representative examples of this class of battery. It is important to note that none of the selected batteries were designed for use under the conditions of these tests. Also their selection for these experiments does not imply a recommendation for their use for any application. The performance of these batteries is being determined to evaluate gelled electrolyte batteries only in a generic sense as compared to conventional liquid electrolyte batteries. It is not intended to evaluate performance of particular suppliers or to compare performance between suppliers.

The batteries from Globe-Union Inc. (2.6 AH), General Electric Co. (2.5 AH) and Gates Energy Products Inc. (2.5 AH) are shown in Figures 1, 2 and 3 respectively. As can be seen, the Globe battery is one integral unit while the others are a package containing three separable 2V cells.

One battery from each vendor was dissected to identify appropriate methods to provide pressure compensation. Small holes were drilled in the battery tops as shown in Figures 1-3 to allow complete flooding of the cells with compensating oil.

Five 6V batteries from each vendor were connected in series during the tests. These fifteen batteries were charge/discharge cycled at 1 atmosphere and approximately $70^{\circ} F$. The first discharge was for $2^{\frac{1}{2}}$ hours at 450 mA. This constituted about a 50% of rated capacity discharge. The second discharge was to a low battery cutoff voltage of 5.0V. As each battery reached cutoff voltage it was removed from the circuit. This pattern was repeated through six discharges (cycle 1-6) until the batteries had received three 50% discharges and three 100% discharges.

Each charge was started at the 250 mA rate and continued until a battery reached 7.5V. That battery was then removed from charge until all the batteries reached 7.5V at 250 mA. Then the batteries were placed back in the circuit at 100 mA and the current was continually tapered to keep the voltage of all cells at or below 7.5V until the current had been reduced to 30 mA. As each battery reached 7.5V at 30 mA it was removed and the taper charge continued until all batteries reached 7.5V at 30 mA. This charging procedure remained the same throughout the entire test series.

The capacities of the five batteries from each manufacturer were then checked. The highest capacity and the lowest capacity batteries from each manufacturer were arbitrarily selected as control batteries. The three middle capacity batteries from each manufacturer became the test batteries. The control batteries and test batteries were cycled again as a series connected assembly of 15 batteries. However, the six control batteries were operated at 1 atmosphere and approximately 70°F, while the test batteries were subjected to a variety of deep ocean conditions described below.

The most widely used fluid for compensating deep ocean batteries is white mineral oil (e.g. Marcol 70). This fluid was judged to be compatible with the construction of the gel cells and the gelled electrolyte. It was, therefore, chosen as the compensating medium for the remainder of these tests.

To determine effects of introducing mineral oil within the cells, six more cycles (7-12) were conducted in the same manner as the first six cycles except all of the test batteries were submerged in mineral oil at 1 atmosphere and 70°F . The control batteries were cycled in air.

To determine effects of pressure upon battery performance the test batteries were next cycled four times (13-16) while submerged in mineral oil at 10,000 psig and about 70° F. The pressure remained at 10,000 psig during charges and discharges.

Partial discharges, by definition, do not provide a measurement of battery capacity. Therefore, in order to increase the number of data points generated, all discharges while operating in the pressure vessel (beginning with cycle 13) were to low battery cutoff voltage for each battery. The control batteries were cycled simultaneously in air.

During the final four cycles (cycles 17-20)the test batteries were charged and discharged in oil at 10,000 psig and 32-35°F. The control batteries were still at 1 atmosphere and 70°F in air. Prior to the second discharge at low temperature (cycle 18) the test batteries were allowed to stand about 62 hours fully charged at 10,000 psig and 32°F. During the other five cycles the stands between the end of charge and start of discharge were less than 16 hours.

TEST RESULTS

THE RESERVE AND ASSESSMENT OF THE PROPERTY OF

The average capacities discharged from the control batteries and test batteries for each cycle are shown in Figures 4-6. (Data from individual batteries are shown in Appendix A.) As previously noted, cycles 1, 3, 5, 7, 9, and 11 were partial discharges. Since full battery capacity was not measured these cycles were not used as data points for the following aralysis. At the end of the sixth cycle the highest and lowest capacity batteries from each vendor were chosen to be control batteries. The other three were designated as test batteries.

The mean capacity for the test batteries from each supplier was the average of the full discharge capacities for all three test batteries during all three full discharges conducted from cycles 1-6 (nine data points). Similarly, the mean of the two control batteries was calculated for each supplier's batteries from the three complete discharges conducted from cycles 1-6 (six data points). A statistical check (using Student t distribution) showed there was no significant difference between the mean capacities (at 1 atmosphere and 70 F) of the control (M_C) and test batteries (M_T) from General Electric and Gates at a significance level of 0.1. That is, the null hypothesis that M_T- M_C= 0 was found to be true at that significance level. However, the batteries chosen from Globe for control batteries had a mean capacity significantly higher than the test batteries, and thus the null hypothesis was rejected. A more complete description of the statistical analysis is given in Appendix B. The mean capacities are shown for each vendor in Table 1 below and also in cycles 1-6 of Figures 4, 5 and 6.

Table 1.

Mean Battery Capacities (AH) for Cycles 1-6
(1 atmosphere, 70°F, in air)

	<u>Globe</u>	General Electric	Gates
Control Batteries ($M_{\mathbb{C}}$)	1.58	1.78	2.08
Test Batteries (M_T)	1.47	1.79	2.11
MT- IAC	-0.11	+.01	+.03
Null Hypothesis: M_{T} - M_{C} = 0	reject	accept	accept

In order to evaluate effects on battery capacity caused by the compensating oil, the mean capacities of the test batteries and the control batteries for cycles 7-12 were compared for changes from the previous six cycles. Any bias existing between the control batteries and test batteries during cycles 1-6 (e.g. the Globe batteries as previously discussed) was subtracted so that only significant changes in the relative mean capacities were evaluated. It was hypothesized that introducing the oil would have no significant effect, that is, after adjusting for bias, M_T - M_C = 0 at a significance level of 0.1. A summary of this analysis is shown in Table 2.

Table 2.

Mean Battery Capacities for Cycles 7-12
(1 atmosphere, 70°F, test batteries in mineral oil)

	<u>Globe</u>	General Electric	Gates
M _C	1.48	1.22	1.77
M_T	1.42	1.49	2.00
$\Delta(M_T - M_C)$	0.05	0.27	0.20
Null Hypothesis: $\Delta(M_T - M_C) = 0$	accept	reject	reject

There was no significant change in the Globe batteries after the introduction of mineral oil, although there was a slight decrease in the magnitude of the difference between the test battery capacities and the control battery capacities (MT- MC). While this indicates a slight relative improvement with the addition of oil, it was not statistically significant. However, the test batteries from GE and Gates showed significant relative battery capacity improvements when operating in oil. Thus, the hypothesis that the oil had no effect was rejected for batteries from these two suppliers. (The compensating oil actually had a positive effect upon battery capacity.)

The test batteries were operated during cycles 13-16 at 10,000 psig submerged in mineral oil. Each battery was discharged to low voltage cutoff on every cycle (no partial discharges). Once again after subtracting the bias introduced by prior testing (cycles 7-12), relative changes in the mean capacities of the control batteries and test batteries were analyzed to determine pressure effects. (As before the hypothesis was no change in capacity would be caused by operating at pressure.) Table 3 shows the results of this analysis.

Table 3.

Mean Battery Capacities for Cycles 13-16 (10,000 psig, 70°F, test batteries in mineral oil)

	<u>Globe</u>	General Electric	<u>Gates</u>
MC	1.56	1.13	1.60
^{M}T	1.49	1.37	1.93
$\Delta(M_T - M_C)$	-0.01	-0.03	0.10
Null Hypothesis: $\Delta(M_T - M_C) = 0$	accept	accept	accept

Thus, there was no statistically significant effect, at the 0.1 significance level, upon mean battery capacity caused by operating at 10,000 psig.

The final test performed involved cooling the pressure vessel and test batteries down to 32°F and cycling them cold at 10,000 psig. The same statistical tests were performed to determine the validity of the hypothesis that no significant relative change in the mean battery capacities would occur due to the cold ambient conditions. Table 4 shows the results of these tests.

Table 4.

Mean Battery Capacities (AH) for Cycles 17-20 (10,000 psig, 32°F, in mineral oil)

	<u>Globe</u>	General Electric	<u>Gates</u>
^M C	1.49	0.88	1.36
M _T	1.23	1.06	1.46
Δ(M _T - M _C)	-0.19	-0.06	-0.23
Null Hypothesis: $\Delta(M_T - M_C) = 0$	reject	accept	reject

By the same type of statistical analysis previously used, the hypothesis that low temperature did not affect battery capacity significantly was tested. For the General Electric batteries the hypothesis was accepted. Although there was some relative drop in the test batteries' capacities, it was not significant at the 0.1 level. However, the hypothesis was rejected for the Globe batteries and the Gates batteries as they showed significant capacity losses when operated cold. Even so, the capacity losses were still less than those previously measured with standard batteries. (Reference (1))

Prior to discharge #18 the test batteries stood cold at 10,000 psig for 62 hours. The average drop in capacity between the first discharge cold (#17) and the second discharge cold (#18), which was preceded by the long stand, was only 0.1 AH. Thus, the long stand while cold did not have a significant effect.

One other comment on the test data concerns the variation of battery performance between manufacturers. When Figures 4-6 are compared, it can be seen that the batteries aged in somewhat different manners. The Globe batteries started with the lowest initial capacity, but degraded very little throughout the test. The Globe batteries were among the highest in capacity at the end of the test. The General Electric batteries had good initial capacities which declined rapidly in early cycling and continued to drop throughout the tests. The Gates batteries had the highest initial capacities and maintained the highest capacities throughout most of the tests. However, the Gates batteries exhibited a greater than average loss of capacity (16%) during low temperature operations. Under these test conditions none of the batteries delivered their rated capacity of 2.5-2.6 AH. However, they were being discharged at approximately the 5-hour rate as opposed to the 20-hour rate upon which the nominal battery capacity ratings are based.

Previous data (reference (1)) concerning similar testing done with liquid electrolyte batteries indicated large capacity losses when the batteries were cycled at low temperature and high pressure. In those tests an average capacity degradation of 35% was noted. Comparing the gelled electrolyte test and control batteries (Table 4) shows that two of the three sets of test batteries were actually outperforming the control batteries even though the test batteries were operating at pressure and cold. This may be due to the small sample size (i.e., the test cells may have just been better cells and the small sample did not allow the cell quality variation to average out). However, if the mean capacity of test batteries, shown in Table 4, is compared with these capacities while warm, shown in Table 3, it can be seen that the test batteries experienced a noticeable drop in capacity when the *emperature was reduced. When this drop in capacity is adjusted for normal capacity decline due to aging (estimated from control batteries) it represents about a 10% capacity degradation. Similar testing of standard lead-acid cells (reference (1)) indicated a capacity loss of about 35% occurred with the battery operated at 6000 psig and 33°F. This capacity degradation increased to a dramatic 70% loss when the batteries were allowed to stand charged for five days prior to discharge. Thus although, the gelled electrolyte lead-acid batteries are adversely affected when operating at reduced temperatures, the capacity loss is considerably less than for standard cells operating in the same environment.

CONCLUSIONS

The second of th

Although the test data resulted from a statistically small sample, the general consistency of results for cells from different vendors and the reaffirmation of lead-acid battery historical effects (e.g. neither mineral oil nor pressure affect their capacity) supports the

conclusions drawn here. Thus, concerning the suitability of lead-acid gel cells for deep ocean applications it is concluded that:

- 1. Commercially available lead-acid gelled electrolyte batteries can be modified to compensate them internally for operation at high pressure.
- 2. White mineral oil provides a suitable compensating fluid with no adverse effects upon battery performance. In fact, some batteries may exhibit slightly improved performance when submerged in mineral oil.
- 3. Operating at high ambient pressures (10,000 psig) has no significant effect upon battery performance.
- 4. Operating at reduced ambient temperatures $(32^{\circ}F)$ and high pressure simultaneously can cause a reduction in available battery capacity.
- 5. The capacity drop for the gelled electrolyte batteries when operated cold and at pressure is considerably less than the loss experienced with standard cells. (10% capacity loss as compared to 35% loss for these tests.)
- 6. Long stands (62 hours as compared to 16 hours) at reduced temperatures do not cause a further reduction in battery capacity for gelled electrolyte batteries.
- 7. Under the special test conditions described in this report, the performance characteristics of gelled electrolyte lead-acid batteries varied appreciably between manufacturers.

ACKNOWLEDGEMENTS

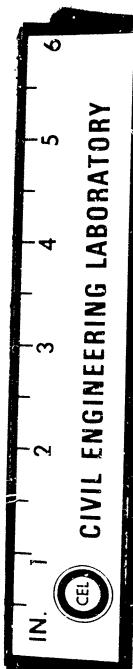
The assistance of Mr. L.W. Tucker and Mr. F.J. Potter was invaluable in conducting these tests and in analyzing the data.

REFERENCE

The house of the state of the s

1. Unpublished data from CEL tests, 1972.





Globe Battery. Figure 1.

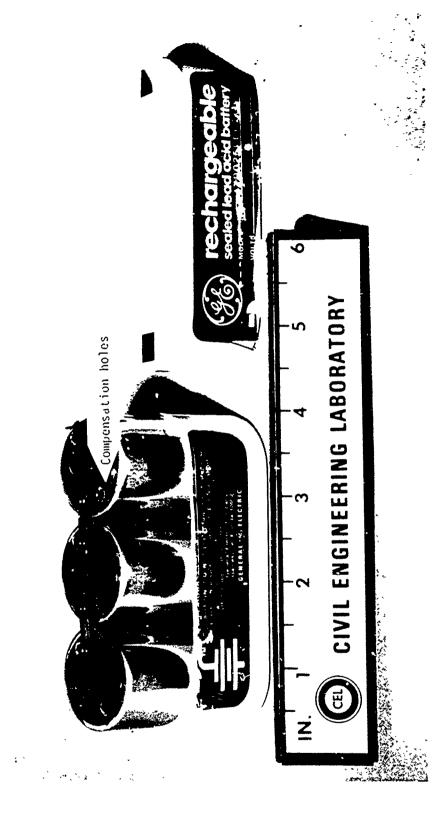


Figure 2. General Blectric Battery.

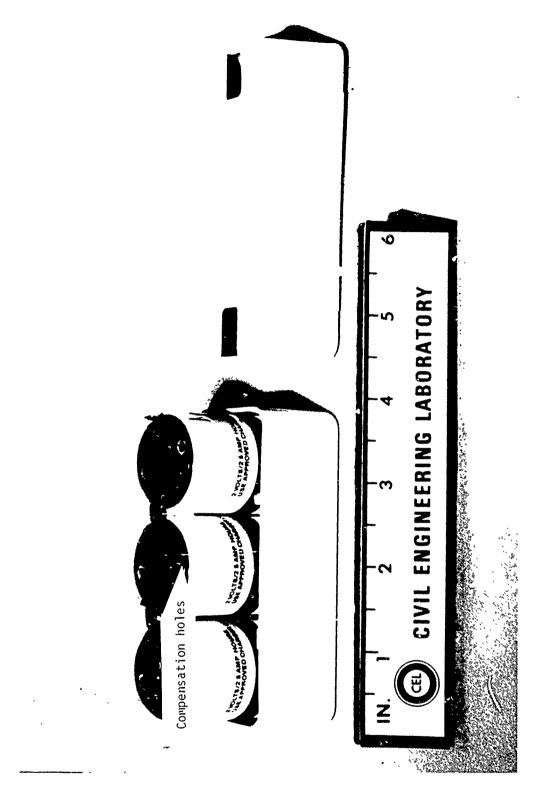
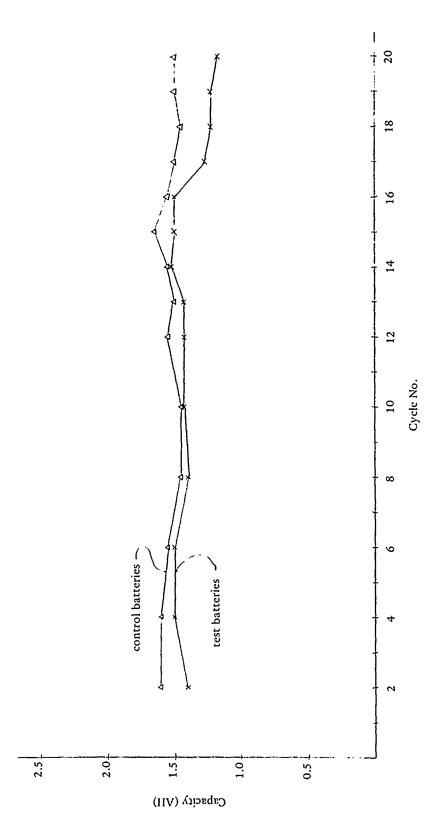
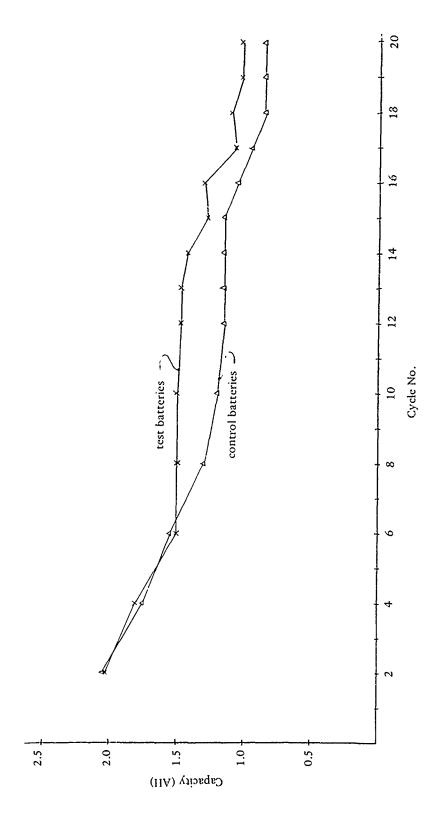


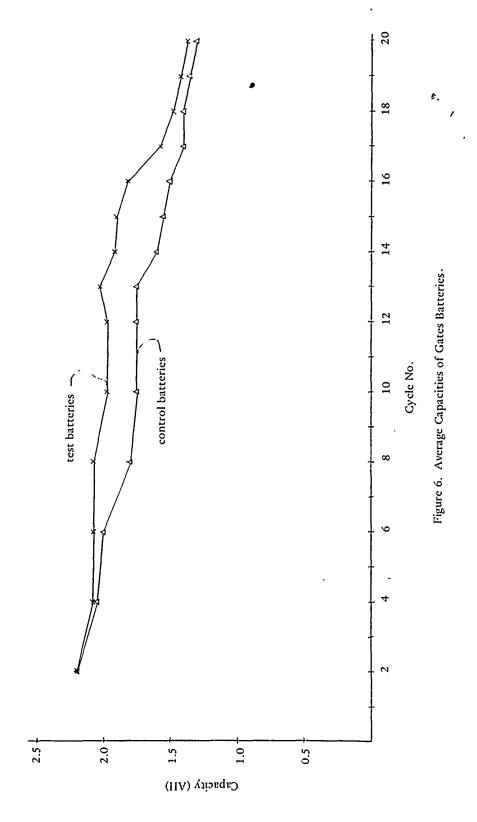
Figure 3. Gates Battery.





Section of the sectio

Figure 5. Average Capacitors of General Electric Batteries.



Appendix A. Individual Battery Capacities.

Test Environment	<u> </u>		×	Air, latm,	atm,	70°F		Œ	Mineral	011,	0il, 1 atm, 70 ⁰ F	۱, 70°	ĮL.	Mine 10,0	Mineral 0il, 10,000 psig,		700F	Mineral Oil, 10,000 psig,	al Oi O psie	1, 32°F	ιι		
cell Type	Cycle No.		2	*	4	*\$	9	*	8	*	2	*=	12	13	14	15	16	17	18	19	. 02	Total	
	0	1.2	1.4	0:	1.5	<u>-</u>	1.5	=	1.4	0.1	1.4	0.	1.4	1.4	1.5	1.5	1.5	1.2	1.2 1.2		1.2	24.4	Test Battery
	_	1.2	1.4	1:0	1.5	0	1.5	-	1.4	0:	1.5	1.0	1.5	1.5	1.6	1.5	1.5	1.3	1,3	1.3	1.2	26.3	Test
Globe	2	0.9	1.4	0	1.5	0:	1.5	=	1.4	0:-	1.4	0	1.4	1.4	1.5	1.5	1.5	1.3	1.2	1.2	1.1	25.3	Test
	8	1.2	1.7	0.1	1.5	0:	1.4	-:	1.3	0	1.3	1.0	1.4	1.3	1.4	1.5	1.4	1.3	1.3	1.4	١.4	25.9	Control
	4	1.2	1.5	<u>.</u>	1.7	- 0:	1.7	Ξ	1.6	0.7	9.	1.0	1.7	1.7	1.7	1.8	1.7	1.7	1.6	9.1	1.6	29.5	Control
	5	1.2	2.2	- 0:	1.9	1.0	1.7	=	1.4	0.1	1.3	0.1	1.2	1.2	1.2	1.2		0.1	6.0	6.0	6.0	24.4	Control
•	ِ و	1.2	2.1	<u>-</u>	1.8	0.7	1.6	=	1.5	0.7	1.5	0:	1.5	1.5	1.5	1.3:1.5	 - 	-	1.3	1.2	1.2	26.9	Test
Electric	7	1.2	2.0	0:	- 8:	1.0	1.4	=	3.5	1.0	1.5	0.1	1.4	1.4	4.	1.2	1.2	0.0	1.0 0.9		6.0	24.9	Test
	_ ∞	1.2	2.0	<u> -</u>	- 8:	<u>-</u>	1.5	=	1.5	0.7	1.5	0.	1.5	1.5	1.4	1.3 1.2		1.1	1.0 1.0		1.0	9:52	Test
	6	1.2	1.9	1.0	9.	<u>-</u>	4.	=	1.2	1.0	-:	1.1 11.0 11.1		7	7	<u>-</u>	1.0	6.0	8.0	8.0	8.0	22.2	Control
	10	1.2	2.1	1.0	1.9	0:	8	=	1.6	1.0	1.6	0:	1.6	1.6	1.5	1.5	1.4	1.3	1.3	1.3	1.3	28.1	Control
	=	1.2	2.3	<u>:</u>	2.2	<u>-</u>	2.2	=	2.0	0.7	6.1	0	1.9	1.9	1.7	1.6	1.6	1.5	1.5	1.4	1.3	31.3	Control
Gates	12	1.2	2.1	0:1	2.0	0:	2.0	=	2.0	1.0	1.9	0.0	1.9	2.0	1.9	1.8	1.8	1.5	1.4	1.4	1.3	31.3	Test
	<u> </u>	13 1.2	2.2	<u>.</u>	2.1	0:	2.1	=	2.1	1.0	2.0	0.:	2.0	2.1	2.0	1.9	1.9	1.6	1.5	1.5	1.4	32.7	Test
ŭ.	14	1.2	2.3	0.1	2.1	0.1	2.1	=	2.1	1.0	2.0	0.	2.0	2.0	1.9	2.0	1.8	9.1	1.5	1.4	1.4	32.5	Test
* Indicates partial discharge	parti	वा का	schar	ge							}												

To analyze the data generated during tests of lead-acid batteries with gelled electrolyte, a methodology of statistical analysis was utilized. This methodology was intended to determine the effects of introduction of compensating fluid, operation at pressure and operation cold upon battery capacity through the use of standard statistical techniques.

The batteries (five from each of three different suppliers) first underwent a break-in period in air. As described in the report the batteries from each supplier were separated into three test batteries (exposed to the test environment) and two control batteries (operated in air). Furthermore, the batteries from each supplier were analyzed separately. The mean capacity for each group in air was estimated from the average capacities of the batteries during the first six cycles (three full discharges). Thus, the means were estimated as follows:

 M_{T} = estimate of the mean of the test battery capacities

$$M_{T} = \frac{J^{\frac{3}{2}} \quad i^{\frac{K}{2}}}{3K} \quad Xj,i$$

and M_{C} = estimate of the mean of the control battery capacities

$$M_{C} = \frac{2}{j^{\frac{\Sigma}{2}} 1} \quad \frac{K}{i^{\frac{\Sigma}{2}} 1} \quad \frac{Yj,i}{2K}$$

where K = number of complete discharges during the test

Xj,i = the capacity measured for the jth test battery during the
 ith complete discharge

The standard deviation of each group of data was also estimated as follows:

 S_T = estimate of the standard deviation of the test battery capacities

$$S_{T} = \sqrt{\frac{\frac{3}{2}\sum_{i=1}^{K} (X_{j}, i - M_{T})^{2}}{N_{T}-1}}$$

and S_C = estimate of the standard deviation of the control battery capacities

$$S_{C} = \sqrt{\frac{j^{\frac{2}{2}} + i^{\frac{K}{2}} + (Yj, i - M_{C})^{2}}{N_{C} - 1}}$$

where N_T = number of data points from the test batteries = 3K

 N_C = number of data points from the control batteries = 2K

The data was then analyzed to determine if a significant difference existed between the test group and the control group, and to determine any bias introduced by the arbitrary division of the batteries into two groups. It was hypothesized that there was no existing bias between the groups and their means would be essentially the same (i.e. M_T - M_C = 0).

Since the sample was small and the true standard deviation was not known, the Student t distribution was chosen as the most appropriate statistic with which to analyze the data. The standard error of the difference between two sample means can be estimated as follows according to reference (1):

 S_D = estimate of the standard error of M_T - M_C

$$S_{D} = \sqrt{\frac{N_{T}S_{T}^{2} + N_{C}S_{C}^{2}}{N_{T}^{+} N_{C}^{-2}}} \qquad \sqrt{\frac{N_{T}^{+} N_{C}}{N_{T}^{-} N_{C}}}$$

Thus using the data and the t statistic with NT+ NC-2 degrees of freedom the difference between MT and MC can be checked for significance. The level of significance α was arbitrarily chosen to be 0.1 for this analysis. This means that the probability of accepting the "null hypothesis", MT- MC = 0, if it is in fact true, is $1-\alpha$ = 0.90. This may be restated as the probability of rejecting the hypothesis when it is in fact true is α = 0.1. From tables in reference (1) the value of the t statistic

can be found (t_{.1,13}=1.771). If the value of the t ratio = $\frac{M_T - M_C}{S_D}$ lies

within the interval from -1.771 to +1.771 then the null hypothesis can be accepted. If the t ratio lies outside this range, then the null hypothesis can be rejected at a significance level of 0.1.

This analysis was conducted for the first test with both groups of batteries operating in air at about 70°F. The values of the various statistics and results of testing for each supplier's batteries is presented in Table B-I.

Table B-I. (All batteries in air at 1 atmosphere and 70°F) (Units in AH)

	<u>Globe</u>	General Electric	<u>Gates</u>
M _T	1.47	1.788	2.11
S _T	0.05	0.24	0.09
M _C	1.58	1.783	2.08
SC	0.13	0.28	0.19
S _C S _D	0.05	0.145	0.08
M _T - M _C	-0.11	.005	0.03
tratio = - SD	-2.20	0.03	0.37
t.1,13	1.771	1.771	1.771
Null Hypothesis: $M_T - M_C = 0$	reject	accept	accept

The second test performed on the batteries was intended to determine any effects caused by allowing the test batteries to flood with mineral oil for the purpose of pressure compensation. Appropriate holes were drilled in the cell tops and the test batteries were submerged in mineral oil during cycles 7-12. The cycling was otherwise identical to cycles 1-6.

When analyzing the data from cycles 7-12 to determine the effects of introducing mineral oil into the cells, it is desirable to "subtract" any bias identified between the means of the test and control group during cycles 1-6. Therefore, prior to applying the test statistic to the difference between the estimated means, M_T- M_C, the difference between the means from the first test, (M_T- M_C)*, was subtracted. Thus, the change in the relative difference between the means was analyzed to determine possible effects of introducing mineral oil. The null hypothesis for this test was $\Delta(M_T-M_C)=(M_T-M_C)-(M_T-M_C)^*=0$. The results of this analysis are shown in Table B-II.

Table B-II. (Test batteries submerged in oil at 1 atmosphere, 70°F) (Units are AH)

M	Globe	General Electric	Gates
^M T	1.42	1.49	2.0
S _T	0.044	0.633	0.071
^M C	1.48	1.22	1.77
S _C	.172	0.117	0.186
S _D	.065	0.044	0.079
M _T - M _C	06	0.27	0.23
$\Delta(M_{T}-M_{C}) = (M_{T}-M_{C}) - (M_{T}-M_{C})*$.05	0.265	0.20
^(M = M)			
$t_{ratio} = \frac{\Delta(M_T - M_C)}{S_D}$	0.769	5.98	2.70
t.1,13	1.771	1.771	1.771
Null Hypothesis:			
$\Delta(M_T - M_C) = 20$	accept	reject	reject

During the third test conducted the test batteries were operated at an ambient pressure of 10,000 psig. Once again, it is desirable to analyze the changes in capacity caused by operating at pressure. Therefore the difference in the mean capacities during the previous test was subtracted from the difference estimated from the data of cycles 13-16. With bias introduced by operation to this point eliminated, the statistical analysis was conducted and the results are shown in Table B-III.

Table B-III (Test batteries submerged in oil at 10,000 psig, 70^{0} F) (Units are AH)

M _T	<u>Globe</u> 1.49	General Electric 1.37	Gates 1.92
S _T	0.051	0.123	0.097
^M C	1.56	1.12	1.60
s_{C}	0.185	0.071	0.151
S_{D}	0.019	0.060	0.058
M _T - M _C	07	0.25	0.33
$\Delta(M_T - M_C)$	01	-0.035	.095
$t_{ratio} = \frac{\Delta(M_T - M_C)}{S_D}$	-0.53	-0.42	1.63
t.1,18	1.734	1.734	1.734
Null Hypothesis: $\Delta(M_T - M_C) = 0$	accept	accept	accept

The final test (cycles 17-20) was designed to determine the effect of low ambient temperature (32°F) in combination with high ambient pressure (10,000 psig) upon battery capacity. In order to look just at the effects of this test the differences between the mean capacities of the test batteries and the control batteries from the previous test (cycles 13-16) were once again subtracted. The statistical analysis of the change in the difference of the sample means was repeated. The results are shown in Table B-IV.

Table IV. (Test batteries submerged in oil at 10,000 psig, $32^{0}\mathrm{F}$) (Units are AH)

	Globe	General Electric	<u>Gates</u>
^{M}T	1.22	1.06	1.46
s _T	0.062	0.124	0.090
•	1.49	0.88	1.36
^M C S _C S _D	0.155	0.071	0.092
Sn	0.053	0.051	0.044
M _T - M _C	-0.26	0.18	0.09
$\Delta(M_T - M_C)$	-0.19	-0.06	-0.23
$t_{\text{ratio}} = \frac{\Delta(M_{\text{T}} - M_{\text{C}})}{S_{\text{D}}}$	-3.673	-1.217	-5.279
t.1,18	1.734	1.734	1.734
Null Hypothesis: $\Delta(M_T - M_C) = 0$	reject	accept	reject

Reference (1): Statistics: Methods and Analysis by Lincoln L. Chao, McGraw-Hill, 1969.

DISTRIBUTION LIST

AFB CESCH, Wright-Patterson; MAC/DET (Col. P. Thompson) Scott, IL; Stinfo Library, Offutt NE

ARCTICSUBLAB Code 54T, San Diego, CA

ARMY BMDSC-RE (H. McClellan) Huntsville AL; ERADCOM Tech Supp Dir. (DELSD-L) Ft. Monmouth, NJ

ARMY COASTAL ENGR RSCH CEN Fort Belvoir VA

ARMY CORPS OF ENGINEERS MRD-Eng. Div., Omaha NE; Seattle Dist. Library, Seattle WA

ARMY ENVIRON. HYGIENE AGCY Water Qual Div (Doner), Aberdeen Prov Ground, MD

ASST SECRETARY OF THE NAVY Spec. Assist Energy (P. Waterman), Washington DC; Spec. Assist Submarines, Washington DC

CINCLANT Civil Engr. Supp. Plans. Ofr Norfolk, VA

CNO Code NOP-964, Washington DC; Code OPNAV 09B24 (H), Code OPNAV 22, Wash DC; Code OPNAV 23, Wash DC; OP987J (J. Boosman), Pentagon

COMFLEACT, OKINAWA PWO, Kadena, Okinawa

COMOCEANSYSPAC SCE, Pearl Harbor HI

DEFENSE DOCUMENTATION CTR Alexandria, VA

DOE Dr. Cohen

DTNSRDC Code 522 (Library), Annapolis MD

MCAS Facil, Engr. Div. Cherry Point NC; J. Taylor, Iwakum Japan

MCRD PWO, San Diego Ca

NAF PWO, Atsugi Japan

NAS Code 18700, Brunswick ME; Dir. Util. Div., Bermuda, ENS Buchholz, Pensacola, FL; PWD Maint, Div., New Orleans, Belle Chasse LA; PWO Belle Chasse, LA; PWO Key West FL; PWO, Glenview IL; SCE Norfolk, VA

NATL RESEARCH COUNCIL Naval Studies Board, Washington DC

NAVACT PWO, London UK

NAVAEROSPREGMEDCEN SCE, Pensacola FL

NAVAL FACILITY PWO, Barbados; PWO, Centerville Bch, Ferndale CA; PWO, Guam

NAVCOASTSYSLAB Code 715 (J. Mittleman) Panama City, FL; Code 715 (J. Quirk) Panama City, FL; Library Panama City, FL

NAVCOMMAREAMSTRSTA SCE Unit I Naples Italy

NAVCOMMSTA Code 401 Nea Makri, Greece: PWO, Exmouth, Australia

NAVCOMMUNIT Cutler/E. Machias ME (PW Gen. For.)

NAVEDTRAPRODEVCEN Tech. Library

NAVELEXSYSCOM Code PME-124-61, Washington DC

NAVENVIRHLTHCEN CO, Cincinnati, OH

NAVEODFAC Code 605, Indian Head MD

NAVFACENGCOld Code 043 Alexandria, VA; Code 044 Alexandria, VA; Code 0451 Alexandria, VA; Code 0454B Alexandria, VA; Code 04B5 Alexandria, VA; Code 101 Alexandria, VA; Code 1023 (T. Stevens) Alexandria, VA; Morrison Yap, Caroline Is.: PC-22 (E. Spencer) Alexandria, VA; PL-2 Ponce P.R. Alexandria, VA

NAVFACENGCOM - CHES DIV. Code FPO-1 (C. Bodey) Wash, DC; Code FPO-1 (Ottsen) Wash, DC; Code FPO-1P1 (Gregory); Code FPO-1SP (Dr. Lewis) Wash, DC; Code FPO-1SP13 (T F Sullivan) Wash, DC; Code FPO-IP12 (Mr. Scola), Washington DC

NAV FACENGCOM - LANT DIV.; Eur. BR Deputy Dir, Naples Italy; RDT&ELO 09P2, Norfolk VA

NAVFACENGCOM - NORTH DIV. (Boretsky) Philadelphia, PA; CO; Code 1028, RDT&ELO, Philadelphia PA; ROICC, Contracts, Crane IN

NAVFACENGCOM - PAC DIV. Code 09DG (Donovan), Pearl Harbor, HI

NAVFACENGCOM - SOUTH DIV. Code 90, RDT&ELO, Charleston SC

NAVFACENGCOM - WEST DIV, Code 04B; RDT&ELO Code 2011 San Bruno, CA

NAVFACENGCOM CONTRACT Eng Div dir, Southwest Pac, Manila, PI; OICC, Southwest Pac, Manila, PI; ROICC Off Point Mugu, CA

NAVNUPWRU MUSE JET Code NPU-30 Port Hueneme, CA

NAVOCEANO Code 1600 Bay St. Louis, MS; Code 3408 (J. Kravitz) Bay St. Louis

NAVOCEANSYSCEN Code 2010 San Diego, CA; Code 4473 Bavside Library, San Diego, CA, Code 52 (H. Talkington) San Diego CA; Code 5204 (J. Stachiw), San Diego, CA, Code 5214 (H. Wheeler), San Diego CA; Code 5224 (R.Jones) San Diego CA; Code 6565 (Tech. Lib.), San Diego CA

NAVPGSCOL D. Leipper, Monterey CA; E. Thornton, Monterey CA

NAVPHIBASE CO, ACB 2 No. lolk, VA, Code S3T, Norfolk VA; Harbor Clearance Unit Two, Little Creek, VA;

OIC, UCT ONE Norfolk, Va

NAVREGMEDCEN SCE (D. Kaye); SCE, Guam

NAVSEASYSCOM Code OOC (LT R. MacDougal), Washington DC: Code SEA OOC Washington, DC

NAVSEC Code 6034 (Library), Washington DC

NAVSHIPYD; Code 202.4, Long Beach CA; Code 202.5 (Library) Puget Sound, Bremerton WA; Code 440 Portsmouth NH; Code 440, Puget Sound, Bremerton WA; Tech Library, Vallejo, CA

NAVSTA Maint, Cont. Div., Guantanamo Bay Cuba; PWO, Mayport FL; Utilities Engr Off. (LTJG A.S. Ritchie), Rota Smin

NAVSTA BISHOPS POINT Harbor Clear. Unit one, Pearl Harbor, H!

NAVSUBSCOL LT J.A. Nelson Groton, CT

NAVSUPPACT Code 413, Seattle WA; LTJG McGarrah, Valleje CA

NAVSURFWPNCEN PWO, White Oak, Silver Spring, MD

NAV FECHTRACEN SCE, Pensacola FL

NAVWPNCEN Code 2636 (W. Bonner), China Lake CA

NAVWPNSTA EARLE PW Office (Code 09C1) Yorktown, VA: PWO, Seal Beach CA

NAVWPNSUPPCEN Code 09 (Boennighausen) Crane IN

NAVXDIVINGU LT A.M. Parisi, Panama City FL

NCBC CEL (CAPT N. W. Petersen), Port Hueneme, CA; CEL AOIC Port Hueneme CA; Code 10 Davisville, RI; Code 156, Port Hueneme, CA

NOAA Librarym Rockville, MD

NORDA Code 410 Bay St. Louis, MS; Code 440 (Ocean Rsch Off, Bay St. Louis MS

NRL Code 8441 (R.A. Skop), Washington DC: Rosenthal, Code 8440, Wash. DC

NSD SCE, Subic Bay, R.P.

NAVOCEANSYSCEN Hawaii Lab (D. Moore), Hawaii

NUSC Code 131 New London, CT: Code EA123 (R.S. Munn), New London CT: Code S332, B-80 (J. Wilcox); Code SB 331 (Brown), Newport RI: Code TA131 (G. De la Cruz), New London CT

OCEANAV Mangmt Info Div., Arlington VA

OCEANSYSLANT LT A.R. Giancola, Norfolk VA

ONR CDR Harlett, Boston MA; BROFF, CO Boston MA; Code 221, Arlington VA; Code 481, Arlington VA; Code 481, Bay St. Louis, MS; Code 700F Arlington VA; Dr. A. Laufer, Pasadena CA

PHIBCB 1 P&E, Coronado, CA

PMTC Pat, Counsel, Point Mugu CA

PWC CO Notfolk, VA; CO, Great Lakes IL; Code 116 (LTJG, A. Eckhart) Great Lakes, IL; Code 120C (Library) San Diego, CA; Code 220.1, Norfolk VA; Code 30C (Boettcher) San Diego, CA; Code 400, Pearl Harbor, HI

U.S. MERCHANT MARINE ACADEMY Kings Point, NY (Reprint Custodian)

US DEPT OF INTERIOR Bureau of Land MNGMNT - Code 733 (T.E. Sullivan) Wash, DC

US GEOLOGICAL SURVEY Off, Marine Geology, Piteleki, Kaston VA

USCG (G-ECV) Washington Dc; (G-MP-3/USP/82) Washington Dc; G-EOE-4/61 (T. Dowd), Washington DC

USCG ACADEMY LT N. Stramandi, New London CT

USCG R&D CENTER LTJG R. Dair, Groton CT

USNA Ocean Sys. Eng Dept (Dr. Monney) Annapolis, MD

CALIF. MARITIME ACADEMY Vallejo, CA (Library)

CATHOLIC UNIV. Mech Engr Dept, Prof. Niedzwecki, Wash., DC

DUKE UNIV MEDICAL CENTER B, Muga, Durham NC

FLORIDA ATLANTIC UNIVERSITY BOCA RATON, FL (MC ALLISTER)

LEHIGH UNIVERSITY BETHLEHEM, PA (MARINE GEOTECHNICAL LAB., RICHARDS); Bethlehem PA (Linderman Lib. No.30, Flecksteiner)

LIBRARY OF CONGRESS WASHINGTON, DC (SCIENCES & TECH DIV)

MAINE MARITIME ACADEMY (Wyman) Castine ME; CASTINE, ME (LIBRARY)

MIT Cambridge MA; Cambridge MA (Rm 10-500, Tech, Reports, Engr. Lib.)

NATL ACADEMY OF ENG. ALEXANDRIA. VA (SEARLE, JR.)

OREGON STATE UNIVERSITY (CE Dept Grace) Corvallis, OR; Corvalis OR (School of Oceanography)

PENNSYLVANIA STATE UNIVERSITY STATE COLLEGE, PA (SNYDER)

SCRIPPS INSTITUTE OF OCEANOGRAPHY LA JOLLA, CA (ADAMS)

SOUTHWEST RSCH INST King, San Antonio, TX; R. DeHart, San Antonio TX

STANFORD UNIVERSITY Engr Lib, Stanford CA: STANFORD, CA (DOUGLAS)

TEXAS A&M UNIVERSITY COLLEGE STATION, TX (CE DEPT); College Station TX (CE Dept. Herbich)

UNIVERSITY OF CALIFORNIA BERKELEY, CA (CE DEPT, GERWICK); La Jolla CA (Acq. Dept, Lib. C-075A);

SAN DIEGO, CA, LA JOLLA, CA (SEROCKI)

UNIVERSITY OF DELAWARE Newark, DE (Dept of Civil Engineering, Chesson)

UNIVERSITY OF HAWAII HONOLULU, HI (SCIENCE AND TECH. DIV.)

UNIVERSITY OF ILLINOIS Metz Ref Rm. Urbana IL; URBANA, IL (DAVISSON); URBANA, IL (LIBRARY), URBANA, IL (NEWARK)

UNIVERSITY OF MASSACHUSETTS (Heronemus). Amherst MA CE Dept

UNIVERSITY OF NEBRASKA-LINCOLN Lincoln, NE (Ross Ice Shelf Proj.)

UNIVERSITY OF NEW HAMPSHIRE DURHAM, NH (LAVOIE)

UNIVERSITY OF RHODE ISLAND KINGSTON, RI (PAZIS); Narragansett RI (Pell Marine Sci. Lib.)

UNIVERSITY OF SO. CALIFORNIA Univ So. Calif

UNIVERSITY OF TEXAS Inst. Marine Sci (Library), Port Arkansas TX

UNIVERSITY OF WASHINGTON SEATTLE, WA (APPLIED PHYSICS LAB): SEATTLE, WA (OCEAN ENG RSCH LAB, GRAY): SEATTLE, WA (PACIFIC MARINE ENVIRON, LAB., HALPERN): Seattle WA (E. Linger)

UNIVERSITY OF WISCONSIN Milwaukee WI (Ctr of Great Lakes Studies)

AGBABIAN ASSOC, C. Bugge, El Segundo CA

AMETEK Offshore Res. & Engr Div

ATLANTIC RICHFIELD CO. DALLAS, TX (SMITH)

BECHTEL CORP. SAN FRANCISCO. CA (PHELPS)

BELGIUM HAECON, N.V., Gent

BRITISH EMBASSY Sci. & Tech. Dept. (J. McAuley), Washington DC

BROWN & CALDWELL E M Saunders Walnut Creek, CA

BROWN & ROOT Houston TX (D. Ward)

CANADA Can-Dive Services (English) North Vancouver, Library, Calgary, Alberta; Lockheed Petro, Serv. Ltd., New Westminster B.C.; Lockheed Petrol Srv. Ltd., New Westminster BC; Nova Scotia Rsch Found, Corp. Dartmouth, Nova Scotia

CHEVRON OIL FIELD RESEARCH CO. LA HABRA, CA (BROOKS)

COLUMBIA GULF TRANSMISSION CO. HOUSTON, TX (ENG. LIB.)

CONCRETE TECHNOLOGY CORP. TACOMA, WA (ANDERSON)

DRAVO CORP Pittsburgh PA (Giannino); Pittsburgh PA (Wright)

NORWAY DET NORSKE VERITAS (Library). Oslo

EVALUATION ASSOC. INC KING OF PRUSSIA. PA (FEDELE)

EXXON PRODUCTION RESEARCH CO Houston TX (A. Butler Jr)

FRANCE Roger LaCroix, Paris

GOULD INC. Shady Side MD (Ches. Inst. Div., W. Paul)

GRUMMAN AEROSPACE CORP. Bethpage NY (Tech. Info. Ctr)

MAKAI OCEAN ENGRNG INC. Kailua, HI

LOCKHEED MISSILES & SPACE CO. INC. L. Trimble, Sunnyvale CA: Sunnyvale, CA (Phillips)

MARATHON OIL CO Houston TX (C. Seay)

MOBIL PIPE LINE CO. DALLAS, TX MGR OF ENGR (NOACK)

NORWAY A. Torum, Trondheim; DET NORSKE VERITAS (Roren) Oslo; 1. Foss. Oslo

OCCAN ENGINEERS SAUSALITO, CA (RYNECKI)

OCEAN RESOURCE ENG. INC. HOUSTON, TX (ANDERSON)

OFFSHORE DEVELOPMENT ENG. INC. BERKELEY, CA

PACIFIC MARINE TECHNOLOGY Long Beach, CA (Wagner)

RAYMOND INTERNATIONAL INC. E Colle Soil Tech Dept, Pennsauken, NJ

SANDIA LABORATORIES Library Div., Livermore CA

SEATECH CORP. MIAMI, FL (PERONI)

SHELL DEVELOPMENT CO. Houston TX (C. Sellars Jr.)

SHELL OIL CO. Houston TX (R. de Castongrene)

TIDEWATER CONSTR. CO Norfolk V v (Fowler)

TRW SYSTEMS CLEVELAND, OH (ENG. LIB.)

UNITED KINGDOM British Embassy (Info Offr), Washington DC, J. Derrington, London, R. Rudham Oxfordshire, Taylor, Woodrow Constr (Stubbs), Southali, Middlesex

WATT BRIAN ASSOC INC. Houston, TX

WESTINGHOUSE ELECTRIC CORP. Annapolis MD (Oceanic Div Lib, Bryan)

ADAMS, CAPT (RET) Irvine, CA

BULLOCK La Canada

R.F. BESIER Old Saybrook CT T.W. MERMEL Washington DC CEC Donofrio, John L., LT